# (19) World Intellectual Property Organization International Bureau





(43) International Publication Date 15 May 2003 (15.05.2003)

**PCT** 

(10) International Publication Number WO 03/039609 A1

(51) International Patent Classification<sup>7</sup>: 27/42, C25D 13/02, 13/20, A61F 2/30

A61L 27/32,

(21) International Application Number: PCT/GB02/04889

(22) International Filing Date: 28 October 2002 (28.10.2002)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data: 0126467.0

3 November 2001 (03.11.2001) GB

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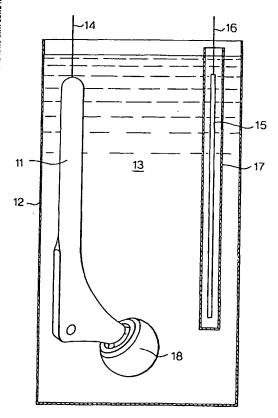
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(81) Designated States (national): A.E., A.G., A.L., A.M., A.T., A.U., A.Z., B.A., B.B., B.G., B.R., B.Y., B.Z., C.A., C.H., C.N., C.O., C.R., C.U., C.Z., D.E., D.K., D.M., D.Z., E.C., E.E., E.S., F.I., G.B., G.D., G.E., G.H., G.M., H.R., H.U., I.D., I.L., I.N., I.S., J.P., K.E., K.G., K.P., K.R., K.Z., L.C., L.K., L.R., L.S., L.T., L.U., L.V., M.A., M.D., M.G., M.K., M.N., M.W., M.Z., N.O., N.Z., O.M., P.H., P.L., P.T., R.O., R.U., S.D., S.E., S.G.,

[Continued on next page]

(54) Title: DEPOSITION OF COATINGS ON SUBSTRATES



(57) Abstract: A two (or more) stage process is described for electrochemically coating a material comprising calcium phosphate onto an electrically conducting substrate, in articular-the electrophoretic deposition of a coating of calcium phosphate, such as hydroxyapatite, on an electrically conducting substrate (such as a bone replacement prosthesis), Surface preparation, precoating and protection precautions are provided to achieve acceptable bond strength for the coating.

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SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

#### Declarations under Rule 4.17:

as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii)) for the following designations AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD,

SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)

— of inventorship (Rule 4.17(iv)) for US only

#### Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

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### Deposition of coatings on substrates

The invention relates to the deposition of calcium phosphate onto a bone replacement prosthesis and more particularly the deposition of calcium phosphate coated vesicles, or droplets having a hydrophobic core coated with calcium phosphate, onto electrically conducting substrates and in particular metallic bone replacement prostheses or parts thereof.

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Patent specification W0 00/00177 describes a vesicle comprising an inner layer which comprises a phospholipid and an outer layer which comprises calcium phosphate. The vesicles may contain pharmaceutically active compounds 15 including compounds which assist the binding of the coating to existing bone (bone growth factors), treat a specific bone disease or any diseased region adjacent to bone, or relieve pain. There is also described a droplet comprising a hydrophobic core, an inner layer which 20 comprises a surfactant, and an outer layer which comprises calcium phosphate. W0 00/00177 also describes a method for producing such vesicles or hydrophobic droplets and a method for depositing such vesicles or droplets on an electrically conducting substrate. 25 method is referred to as electrolytic deposition but is perhaps more accurately described as electrophoretic deposition.

While this deposition method can be carried out at low enough temperatures to avoid degrading the contents of the vesicles or droplets, a problem is achieving satisfactory adhesion of the deposited material to the substrate.

It is an object of the present invention to overcome or ameliorate this problem. It is a further object of

this invention to provide more generally a method of preparing an electrically conducting substrate and depositing a coating thereon so as to achieve good distribution and adhesion of the coating on the substrate.

The invention provides in one of its aspects a method for producing an adherent coating of a coating material comprising calcium phosphate on an electrically conducting substrate, which method comprises first exposing the substrate to an electrochemical surface pretreatment using a solution comprising water for removing contaminants and/or effecting a surface conversion and/or applying a primary coating to form a surface layer or surface compound layer adapted to bond to a subsequently deposited coating, and depositing upon the pre-treated surface the coating material by an electrochemical deposition process from a solution comprising calcium and phosphate ions or a suspension of calcium phosphate 20 particles in liquid.

In a preferred embodiment, the electrochemical surface pre-treatment is a treatment for removing contaminants and/or effecting a surface conversion.

25 Preferably the pre-treatment comprises effecting a surface conversion.

The electrically conducting substrate may be any substrate which conducts electricity or which has a conducting surface. The substrate is preferably metal, more preferably titanium, a titanium alloy or a cobalt/chromium alloy, most preferably a titanium alloy.

The electrochemical surface pre-treatment may

35 comprise electrochemical etching to dissolve a thin

surface layer of the substrate surface. Preferably,

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where the method is applied to a metal substrate, the electrolyte composition and impressed electrical current are selected so as to reveal the micro-structure of the metal by dissolving the grain boundaries at a faster rate and thereby roughen the surface. Mineral acid electrolyte will normally be suitable, with a complexant, such as fluoride, where the substrate metal is titanium. Application of ultrasound and/or current reversal is effective for creating a suitably rough surface.

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The electrochemical surface pre-treatment takes place using a solution comprising water as a solvent. The solution may comprise a mixture of solvents including water. Preferably the solution comprises at least 20% water, more preferably 50% water, more preferably 70% and most preferably 90% or more of water. In a particularly preferred embodiment, an aqueous solution is used and water is the only solvent. In a preferred embodiment the solution is an aqueous phosphate electrolyte.

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During the surface conversion, the surface of the electrically conducting substrate may be converted, for example, into an oxide, a metal salt or a mixed oxide and metal salt to form a layer that is integral with the surface of the substrate. The oxide is typically a hydrous oxide. Where the surface conversion layer is a mixture of oxide and metal salts, both the oxide and the metal salt are derived from the metal of the substrate. The metal salt is typically insoluble in water.

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Where the electrically conducting substrate is titanium or a titanium alloy then the surface conversion may convert the surface of the metal to form an oxide film on which calcium phosphate is deposited in the electrochemical deposition step of the process. The surface conversion layer may also be a mixed oxide and

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titanium salt layer where the titanium salt is one that is insoluble in water. For example, a mixed oxide and phosphate layer can be formed when phosphoric acid is used in the surface conversion process.

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Where the electrically conducting substrate is a cobalt/chromium alloy, the surface conversion may be achieved by dissolving the surface of the substrate in a phosphate medium in order to form metal phosphate salts on the surface of the substrate.

The electrochemical deposition process may comprise electrophoretic deposition of particulate material from a suspension thereof in liquid, or electrocoagulation, or electroprecipitation from a solution of a desired coating material or precursor therefor, or electrolytic deposition from a solution of a desired coating material or precursor therefor.

More than one electrochemical deposition process may be used to build a series of coating layers chosen, for example, to optimise binding efficiency of one layer upon the immediately preceding layer.

25 A pre-treatment surface coating, or the deposited coating, or one or more of the deposited coating layers may be subjected to a plasma fusing action. This is conveniently achieved by allowing the applied voltage to rise as the surface coating or coating layer deposits on the substrate and raises the electrical resistance across the substrate surface, such that plasma forms in pores of the deposited coating and thereby generates heat sufficient to melt and fuse the coating. In this way increased coating density(if wanted) and adhesion to the substrate can be achieved.

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The method according to the invention has particular application to the coating onto a substrate electrically conducting surface of vesicles comprising an inner phospholipid coated with an outer layer of calcium phosphate or of droplets comprising an inner hydrophobic core coated with an outer layer of calcium phosphate.

In one embodiment, prior to the electrophoretic deposition of the calcium phosphate or the said vesicles or droplets, one or more interlayers is provided on the cleaned and roughened metal surface. The first interlayer is the surface conversion layer. For certain metals, such as titanium, the surface conversion coating layer can conveniently be in the form of an adherent oxide film produced by anodic polarisation. Periodic polarity reversal may be applied to increase the thickness of the oxide film layer formed. As an alternative, a phosphate conversion layer can be formed in phosphoric acid.

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For certain metals, such as a cobalt/chromium alloy, it is important to avoid corrosion of the metal surface as this not only will undercut subsequent coatings, but also introduce corrosion products into the coating. The presence of such corrosion products may be undesirable for a bone replacement prosthesis where the corrosion products may subsequently leach into body fluids or tissue. A suitable protective interlayer on cobalt/chromium alloy can be provided by surface dissolution into a neutral/alkaline phosphate solution which creates an adherent insoluble phosphate layer which forms the surface conversion layer. It is typically advantageous to use polarity reversal when forming a surface conversion layer on a cobalt/chromium alloy.

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A graduation of coating properties through the

thickness of the coating may be engineered by appropriate control of coating conditions such as particle size, zeta potential, applied electrical potential, pH, temperature. This may conveniently be achieved by using multiple 5 immersion tanks. For example, a hydroxyapatite coating may be laid down on a surface conversion coating, and on this an outer layer of hydroxyapatite vesicles. The inner layer of hydroxyapatite may be subjected to a plasma treatment as described above to increase its density and 10 adhesion to the substrate. Such an inner layer provides a strongly adherent key onto which hydroxyapatite vesicles (which need to be deposited without significant heating as explained in WO 00/00177) can then be deposited by electrophoresis. This process may also be performed for 15 other forms of calcium phosphate or using other forms of calcium phosphate as the coating layer.

Insoluble Hydroxyapatite is the form of calcium phosphate required when resorption into body fluids is to 20 be avoided. In certain circumstances, it is specifically desired to provide a coating, or part of a coating, which is resorbable into body fluids. In that case conditions of deposition are chosen which deposit calcium phosphate in a different, soluble, form.

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Coating layers grown so as to have columnar pores, with control of the pore diameter and length, may be used to provide a reservoir within the pores for pharmaceutically active compounds, such as antibiotics, 30 pain relief agents, etc.

Control of the fluid viscosity, surface charge and water solubility can provide for progressive release of an active agent diffusing out of the coating.

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The passage of the electrophoretic deposition

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current can sometimes result in the production of gas bubbles at a substrate metal surface, particularly in aqueous media. This is undesirable as bubble growth can disrupt the coating and/or the presence of the bubbles adversely affects the adhesion of the coating. To avoid this, there is preferably included in the deposition suspension a sacrificial component, that is a component which is easily reduced at a cathode or oxidised at an anode (whichever is required) to create a water soluble rather than a gaseous product. An alternative approach is to use liquids which comprise organic solvents, for example alcohols, glycols or ketones.

Calcium phosphate is soluble in acidic media. As 15 described in WO 00/00177 calcium phosphate in solution is caused to crystallise around a vesicle or droplet using ultrasonics to disperse a water immiscible solvent phase containing the dissolved pharmaceutically active agent to create dispersed droplets around which the calcium 20 phosphate forms a coating. We have found that calcium phosphate crystal formation is improved in this process by progressively reducing the solution acidity. action to raise the pH of the solution may be achieved by chemical reaction such as introduction at a controlled 25 rate of formaldehyde into nitric acid, or by electrodialytic action in which anions are replaced with hydroxyls, or by electrolytic action such as nitrate reduction at a cathode. These processes for changing the pH can also aid the deposition of calcium phosphate 30 particles from a suspension.

Normally, the perturbation of pH at the depositing surface encourages the suspended fine particles to coagulate onto the roughened substrate metal surface,

thus creating an adherent layer. For this, it is important to optimise the conditions of the ions of

calcium, phosphate, complexant and the pH of the solution. Further improvement of the bond strength between the deposited particles may be achieved by incorporating soluble resins (polycarboxylic acids) in the solution. These may either dry out to create an increased bond strength, or be cross-linked in-situ through reactions occurring at the substrate surface. The polymeric materials used would need to be biocompatible for implant applications. Organically modified ceramics may also be included as part of the suspension to provide a key onto which the polymer could bind, as well as creating an environment encouraging the formation of a stable structure around the vesicles or droplets.

15

In a further embodiment, the present invention provides a method for producing an adherent coating of a coating material comprising vesicles on an electrically conducting substrate, which method comprises first 20 exposing the substrate to an electrochemical surface pretreatment for removing contaminants and/or effecting a surface conversion and/or applying a primary coating to form a surface layer or surface compound layer adapted to bond to a subsequently deposited coating, and depositing 25 upon the pre-treated surface by an electrochemical deposition process a coating material comprising vesicles comprising an inner phospholipid coated with an outer layer of calcium phosphate. Preferably the electrochemical surface pre-treatment is a treatment for 30 removing contaminants and/or effecting a surface conversion layer or the pre-treatment comprises effecting a surface conversion layer. Preferably the electrochemical deposition process is electrophoretic deposition. The electrochemical pre-treatment may take 35 place in any solvent or solvent mixture, such as an organic solvent including one or more of an alcohol,

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glycol or ketone, or water or a mixture thereof.

The invention includes a substrate when provided with a surface coating by a method as aforesaid.

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A specific example of a method embodying the invention will now be described with reference to the drawing filed herewith, which is a diagrammatic representation of a hip implant prosthesis undergoing treatment in an electrochemical treatment bath.

In this example, hip prosthesis 11 is made of titanium and is shown in an electrochemical bath 12 filled with appropriate treatment solution or electrolyte 13. The prosthesis 11 is connected via lead 14 to an appropriate source of electrical potential. Electrode 15 is connected via lead 16 to the source of electrical potential. There is also provided a membrane 17 encompassing the electrode 15 for use when the electrode 15 is connected to a positive potential and it is required to protect the prosthesis 11 from any ions dissolved into the electrolyte 13 from the electrode 15. The membrane 17 may have to be removed when a negative potential is applied to electrode 15.

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Parts of the prosthesis 11, primarily the ball joint 18, which are not to be coated are provided with a protective insulating layer, for example of tape. For the first operation to remove contaminants from the metal surface of the titanium prosthesis 11, a surface layer of the metal is electrolytically removed using an acid bath containing fluoride with the prosthesis 11 connected to a positive electrical potential supply and the electrode 15 connected to the negative supply. Application of ultrasound to the bath and/or periodic current reversal helps to introduce surface roughness during this process.

With a suitable change in bath conditions and polarity, the second stage creation of an adherent oxide film on the titanium prosthesis 11 takes place. This is the surface conversion coating. If phosphoric acid is present in the bath then a mixed phosphate and oxide surface conversion coating is formed.

The prosthesis 11 is then moved to another bath, or

10 the bath 12 is washed out and its contents replaced, to
provide a solution containing calcium and phosphate ions,
and a potential applied to deposit a layer of
hydroxyapatite on to the clean titanium surface. At this
stage nitrate may be included in the solution as a

15 sacrificial component to prevent the release of gas
bubbles from the titanium surface.

To densify this layer of hydroxyapatite and improve its adhesion to the substrate prosthesis 11, the applied deposition voltage is allowed to rise as the layer thickness increases (and the surface electrical resistance increases correspondingly), such that plasma forms in the pores of deposited hydroxyapatite and generates sufficient heat to cause melt fusing onto the substrate.

In the final step, a treatment suspension 13 is provided which contains vesicles, each of which comprises an inner layer which comprises a phospholipid and an outer layer which comprises calcium phosphate. The vesicles may contain pharmaceutically active compounds including compounds which assist the binding of the coating to existing bone (bone growth factors), treat a specific bone disease or any diseased region adjacent to bone, or relieve pain. A potential is applied to deposit a coating layer of hydroxyapatite vesicles which bond to

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the interlayer of hydroxyapatite applied in the previous stage. During this deposition temperature rise is controlled, ideally to maintain temperature close to ambient so as not to damage constituents of the vesicles.

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In this way adhesion of the coatings with a static shear strength of at least 15 Mpa can be achieved.

The invention is not restricted to the details of

the foregoing example. For instance the method may be
applied to other metal prostheses made, for example of an
alloy of cobalt and chromium. The prosthesis provided
with a simple layer of strongly adherent hydroxyapatite
(ie without vesicles) has usefulness in its own right,

although the strong bonding at low temperature of
hydroxyapatite vesicles is a key characteristic of the
product of the example. The latter can be deposited
directly onto the first conversion layer (titanium oxide)
of the prosthesis, if desired.

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The final stage deposition need not necessarily be of vesicles as described, but may be of droplets having any hydrophobic core encased in calcium phosphate.

Incorporation of other elements by substitution in the calcium phosphate may be desirable. The other elements are preferably alkaline earth metals, more preferably magnesium, strontium or barium or a mixture thereof. For example, substituted magnesium can act to promote bone growth, or substituted strontium can affect adhesion, or the retention or release of a drug contained in the vesicles.

The method has application for other electrically

35 conducting substrates and other coating materials. A

particular advantage of electrophoretic deposition and

other electrochemical deposition processes, used for depositing electrically insulating coatings, is that the deposited layer is self levelling. This is because the resistance of thick regions reduces current flow in those regions and thus encourages greater deposition in regions where the coating is thinner.

Thus, for example, the method has particular value for providing a uniform thickness coating on a

10 corrugated, or similar uneven surface, such as a catalyst coating on a corrugated metal (eg FECRALLOY steel) substrate.

#### Claims

- A method for producing an adherent coating of a coating material comprising calcium phosphate on an
   electrically conducting substrate, which method comprises first exposing the substrate to an electrochemical surface pre-treatment using a solution comprising water for removing contaminants and/or effecting a surface conversion and/or applying a primary coating to form a
   surface layer or surface compound layer adapted to bond to a subsequently deposited coating, and depositing upon the pre-treated surface the coating material by an electrochemical deposition process from a solution comprising calcium and phosphate ions or a suspension of
   calcium phosphate particles in liquid.
- A method as claimed in claim 1, wherein the electrochemical surface pre-treatment comprises electrochemical etching to dissolve a thin surface layer
   of the substrate surface.
- A method as claimed in claim 2, applied to a metal substrate, wherein, the electrolyte composition and impressed electrical current are selected so as to reveal the micro-structure of the metal by dissolving the grain boundaries at a faster rate and thereby roughen the surface.
- 4. A method as claimed in claim 2 or claim 3, wherein 30 ultrasound and/or current reversal is applied to assist the creation of a rough surface on the etched substrate metal surface.
- A method as claimed in any of the preceding claims,
   wherein the electrochemical deposition process comprises electrophoretic deposition of particulate material from a

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suspension thereof in liquid.

- 6. A method as claimed in any of claims 1 to 4, wherein the electrochemical deposition process comprises5 electrocoagulation.
- 7. A method as claimed in any of claims 1 to 4, wherein the electrochemical deposition process comprises electroprecipitation from a solution of a desired coating 10 material or precursor therefor.
- A method as claimed in any of claims 1 to 4, wherein the electrochemical deposition process comprises electrolytic deposition from a solution of a desired
   coating material or precursor therefor.
  - 9. A method as claimed in any of the preceding claims, wherein more than one electrochemical deposition process is used to build a series of coating layers.

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10. A method as claimed in any of the preceding claims, wherein a pre-treatment surface coating, or the deposited coating, or one or more of the deposited coating layers is subjected to a plasma fusing action.

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- 11. A method as claimed in in any of claims 1 to 10, wherein the pre-treatment comprises treating the substrate to remove surface contaminants and the coating comprises an inner phospholipid coated with an outer layer of calcium phosphate or of droplets comprising an
- layer of calcium phosphate or of droplets comprising an inner hydrophobic core coated with an outer layer of calcium phosphate.
- 12. A method as claimed in any of the preceding claims, 35 wherein, prior to the electrophoretic deposition of the coating, one or more interlayers is provided on the

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cleaned and roughened metal surface.

- 13. A method as claimed in claim 12, wherein the substrate metal comprises titanium and the first5 interlayer is a conversion coating layer in the form of an adherent oxide film produced by anodic polarisation.
- 14. A method as claimed in claim 12, wherein the substrate metal comprises titanium and the first10 interlayer is a phosphate conversion layer formed in phosphoric acid.
- 15. A method as claimed in claim 12, wherein the substrate metal comprises a cobalt/chromium alloy and the first interlayer comprises an adherent insoluble phosphate layer provided by surface dissolution into a neutral/alkaline phosphate solution.
- 16. A method as claimed in any of the preceding claims,
  20 wherein a hydroxyapatite coating is laid down on the
  surface conversion coating, and on the hydroxyapatite
  layer an outer layer of hydroxyapatite vesicles is
  applied.
- 25 17. A method as claimed in any of the preceding claims, wherein coating layers are grown so as to have columnar pores to provide a reservoir within the pores for pharmaceutically active compounds.
- 30 18. A method as claimed in any of the preceding claims, wherein there is included in the deposition suspension a sacrificial component to create a water soluble rather than a gaseous product as a result of any electrolytic action at the surface of the substrate metal.
  - 19. A method as claimed in any of the preceding claims,

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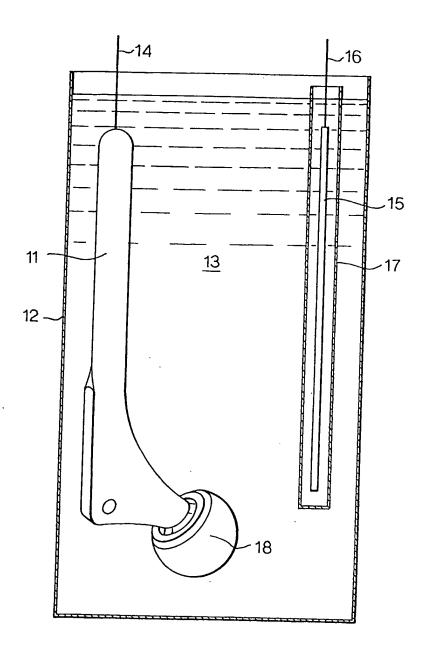
wherein the solution or liquid comprises one or more alcohols, glycols or ketones.

- 20. A method as claimed in any of the preceding claims,5 wherein soluble polycarboxylic acid is incorporated in the solution or liquid.
- 21. A method as claimed in claim 20, wherein organically modified ceramics are included as part of the suspension10 to provide a key onto which the polycarboxylic acid binds.
- 22. A method as claimed in any of the preceding claims wherein the pH of the solution and/or suspension is15 progressively raised by chemical reaction involving introduction at a controlled rate of an appropriate chemical reagent.
- 23. A method as claimed in any of the preceding claims wherein the pH of the solution and/or suspension is progressively raised by electrodialytic action in which anions are replaced with hydroxyls.
- 24. A method as claimed in any of the preceding claims25 wherein the pH of the solution and/or suspension is progressively raised by electrolytic action.
- 25. A method as claimed in any of the preceding claims wherein other elements are incorporated in the calcium30 phosphate coating by substitution.
  - 26. A method as claimed in claim 25 wherein the other elements are magnesium, strontium or barium or a mixture thereof.
  - 27. A method for producing an adherent coating of a

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coating material comprising vesicles on an electrically conducting substrate, which method comprises first exposing the substrate to an electrochemical surface pretreatment for removing contaminants and/or effecting a surface conversion and/or applying a primary coating to form a surface layer or surface compound layer adapted to bond to a subsequently deposited coating, and depositing upon the pre-treated surface by an electrochemical deposition process a coating material comprising vesicles comprising an inner phospholipid coated with an outer layer of calcium phosphate.

28. An electrically conducting substrate when provided with a coating by a method as claimed in any of the preceding claims.



### INTERNATIONAL SEARCH REPORT

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C. DOCUI	MENTS CONSIDERED TO BE RELEVANT			
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